

## IN THE CLAIMS

The listing of claims will replace all prior versions, and listings, of claims in the application:

1. (currently amended) An antimicrobial peptide consisting of the sequence Thr-Ala-Thr-Val-Thr-Val (SEQ ID NO:1 ),  
~~corresponding to active sites of amino terminal extension of subunits assembling surface adhesive organelles of pathogenic Gram-negative bacteria,~~  
~~said peptide further being capable of preventing formation of said surface adhesive organelles by preventing self polymerization of equal peptide units.~~
2. (currently amended) The antimicrobial peptide according to claim 1, wherein the antimicrobial peptide is capable of inhibiting formation of surface adhesive organelles of pathogenic Gram negative bacteria by inhibiting self polymerization of equal peptide units. ~~corresponds to the active sites of amino terminal extension of subunits assembling surface adhesive organelles of *Escherichia coli*.~~
3. (canceled)
4. (canceled)
5. (canceled)
6. (currently amended) The antimicrobial peptide according to claim 2, wherein the antimicrobial peptide ~~prevents~~ inhibits binding of equal protein units with each other and is capable of binding with a binding constant of  $10^3$  M or higher with a polymerizing protein unit.

7. (currently amended) The antimicrobial peptide according to claim 6, wherein the antimicrobial peptide is effective in inhibiting ~~preventing~~ self-polymerization of bacterial virulence organelles in a concentration less than  $10^{-4}$  M.

8. (canceled)

9. (withdrawn ) A method to treat bacterial infections by preventing self –polymerization of equal peptide units of bacterial surface adhesive organelles, thereby preventing formation of said surface adhesive organelles,

said method further comprising administering to a patient a therapeutically active amount of the antimicrobial peptide of claim 1.

10. (withdrawn ) The method according to claim 9, wherein the antimicrobial peptide is further bound to a small molecular or macromolecular substance, thereby increasing the stability of the peptide.

11. (withdrawn) The method according to claim 9, wherein the antimicrobial peptide is applied orally, subcutaneously, or injected into blood circulation.

12. (withdrawn) The method according to claim 11, wherein the antimicrobial peptide is applied in a concentration between  $10^{-4}$  M to  $10^{-10}$  M in sera during prevention or treatment of microbial infections.

13. (withdrawn ) A method for obtaining antimicrobial peptides according to claim 1, the method comprising the steps of:

- a) Cultivating a non pathogenic test microbial strain expressing recombinant self-polymerizing surface organelles of a bacterium;
- b) Adding a candidate compound of antibacterial drug into a mixture of the self-polymerizing organelles in an appropriate concentration;
- c) Investigating degree of polymerization of the surface organelle; and

d) Judging that the compound has an antivirulence action when the polymerization is lowered.

14. (withdrawn) The method of claim 13, wherein the microbial strain expressing recombinant surface organelles is *Escherichia coli* and the polymerizing surface organelle is from *Yersinia*.

15. ( canceled )

16. (canceled)

17. (canceled)